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| (54) Title: PROCESS FOR THE PREPARATION OF N-[(1- ⁿ BUTYL-4-PIPERIDYL)METHYL]-3,4-DIHYDRO-2H-[1,3]OXAZINO[3,2-a]INDOLE-10-CARBOXAMIDE AND SALTS AND INTERMEDIATES IN THE PROCESS | | | | | |
| (57) Abstract | | | | | |
| A process for the preparation of SB 207266 or a pharmaceutically acceptable salt thereof, which process comprises the reaction of N-(1- ⁿ butyl-4-piperidyl)methylamine with a compound of formula (A), wherein R is alkyl. | | | | | |
| <p style="text-align: center;">(A)</p> | | | | | |

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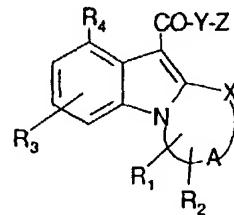
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PROCESS FOR THE PREPARATION OF N-[(1-*n*-BUTYL-4-PIPERIDYL)METHYL]-3,4-DIHYDRO-2H-[1,3] OXAZINO[3,2-a] INDOLE-10-CARBOXAMIDE AND SALTS AND INTERMEDIATES IN THE PROCESS

This invention relates to a new synthetic process to a compound having
5 pharmacological activity.

WO 93/18036 (SmithKline Beecham plc) describes compounds having 5-HT₄ receptor antagonist activity of formula (I), or a pharmaceutically acceptable salt thereof:



10

(I)

wherein

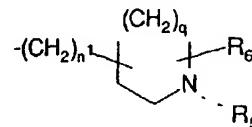
X is O, S, SO₂, CH₂, CH or NR wherein R is hydrogen or C₁₋₆ alkyl;

15 A is a saturated or unsaturated polymethylene chain of 2 - 4 carbon atoms;

R₁ and R₂ are hydrogen or C₁₋₆ alkyl;R₃ is hydrogen, halo, C₁₋₆ alkyl, amino, nitro or C₁₋₆ alkoxy;R₄ is hydrogen, halo, C₁₋₆ alkyl or C₁₋₆ alkoxy;

Y is O or NH;

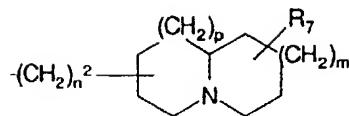
20 Z is of sub-formula (a), (b) or (c):



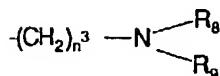
(a)

25

- 2 -



(b)



5

(c)

wherein

n¹ is 1, 2, 3 or 4; n² is 0, 1, 2, 3 or 4; n³ is 2, 3, 4 or 5;

q is 0, 1, 2 or 3; p is 0, 1 or 2; m is 0, 1 or 2;

10 R₅ is hydrogen, C₁₋₁₂ alkyl, aralkyl or R₅ is (CH₂)_z-R₁₀ wherein z is 2 or 3 and R₁₀ is selected from cyano, hydroxyl, C₁₋₆ alkoxy, phenoxy, C(O)C₁₋₆ alkyl, COC₆H₅, -CONR₁₁R₁₂, NR₁₁COR₁₂, SO₂NR₁₁R₁₂ or NR₁₁SO₂R₁₂ wherein R₁₁ and R₁₂ are hydrogen or C₁₋₆ alkyl; and

R₆, R₇ and R₈ are independently hydrogen or C₁₋₆ alkyl; and

15 R₉ is hydrogen or C₁₋₁₀ alkyl;
or a compound of formula (I) wherein the CO-Y linkage is replaced by a heterocyclic bioisostere;

having 5-HT₄ receptor antagonist activity.

20 Examples of alkyl or alkyl containing groups described herein include C₁, C₂, C₃, C₄, C₅, C₆, C₇, C₈, C₉, C₁₀, C₁₁ or C₁₂ branched, straight chained or cyclic alkyl, as appropriate. C₁₋₄ alkyl groups include methyl, ethyl, *n*- and *iso*-propyl, *n*-, *iso*-, *sec*- and *tert*-butyl. Cyclic alkyl includes cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and cyclooctyl.

25 The pharmaceutically acceptable salts of the compounds of the formula (I) include acid addition salts with conventional acids such as hydrochloric, hydrobromic, boric, phosphoric, sulphuric acids and pharmaceutically acceptable organic acids such as acetic, tartaric, maleic, citric, succinic, benzoic, ascorbic, methanesulphonic, α -keto glutaric, α -glycerophosphoric, and glucose-1-phosphoric acids.

Examples of pharmaceutically acceptable salts include quaternary derivatives of the compounds of formula (I) such as the compounds quaternised by compounds $R_X\text{-}T$ wherein R_X is C_{1-6} alkyl, phenyl- C_{1-6} alkyl or C_{5-7} cycloalkyl, and T is a radical corresponding to an anion of an acid. Suitable examples of R_X include methyl, 5 ethyl and *n*- and *iso*-propyl; and benzyl and phenethyl. Suitable examples of T include halide such as chloride, bromide and iodide.

Examples of pharmaceutically acceptable salts also include internal salts such as N-oxides.

The compounds of the formula (I), their pharmaceutically acceptable salts, 10 (including quaternary derivatives and N-oxides) may also form pharmaceutically acceptable solvates, such as hydrates, which are included wherever a compound of formula (I) or a salt thereof is referred to.

Example 3 describes the hydrochloride salt of the compound of formula (I):

15 A is $-\text{CH}_2\text{-(CH}_2\text{)}_r\text{-CH}_2$ - wherein r is 1;
 R_1 and R_2 are hydrogen;
 R_3 is hydrogen;
 R_4 is hydrogen;
 Y is NH ; and
20 Z is of sub-formula (a), and is of structure (i):



This compound is $\text{N-}[(1\text{-}n\text{butyl-4-piperidyl)methyl]-3,4-dihydro-2H-[1,3]oxazino[3,2-a]indole-10-carboxamide}$ SB 207266, (the hydrochloride salt is 25 SB 207266-A) which is being developed by SmithKline Beecham plc as the active ingredient in a medicament for treatment of irritable bowel syndrome.

Example 3 of WO 93/18036 describes a method of preparation of SB 207266-A from $\text{N-}[(1\text{-}n\text{butyl-4-piperidyl)methyl]indole-3-carboxamide}$ (i.e. the compound corresponding to SB 207266, without the oxazino moiety), by reacting 30 with N-chlorosuccinimide and 3-bromo-1-propanol, followed by treatment with

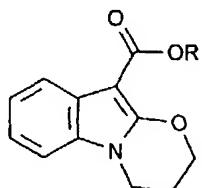
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sodium carbonate. N-[(1-nbutyl-4-piperidyl)methyl]indole-3-carboxamide is prepared by coupling N-(1-nbutyl-4-piperidyl)methylamine with a indole-3-carboxylic acid.

5 An alternative process for preparing SB 207266-A has now been discovered which involves the use of the N-(1-nbutyl-4-piperidyl)methylamine intermediate at a later stage in the process thus resulting in an increased yield of SB 207266-A relative to the amount of this intermediate, which is relatively expensive to produce.

Accordingly, the present invention provides a process for the preparation of SB 207266 or a pharmaceutically acceptable salt thereof, which process comprises the reaction of N-(1-nbutyl-4-piperidyl)methylamine with a compound of formula (A):

10



(A)

wherein R is alkyl, such as methyl or ethyl.

15 The compound of formula (A) wherein R is methyl is methyl 3,4-dihydro-2H-[1,3]-oxazino[3,2-a]indole-10-carboxylate.

The conditions and reagents for this reaction are similar to those described in the literature. A mixture of the amine and ester in a suitable solvent (eg toluene) is treated with a solution of trimethylaluminium in toluene or hexanes at ambient temperature. The resulting solution is then heated, preferably to reflux (112 °C) for 20 about four hours until the reaction is complete. The reaction is cooled to about 70°C and quenched by cautious addition of aqueous sodium hydroxide solution. The aqueous layer is separated and the mixture washed once more with caustic and twice with water, maintaining the temperature at about 70°C. The product is isolated as described in the attached Example.

25 Alternative catalysts include $\text{NaH}_2\text{Et}_2\text{Al}$ which is used in a similar way to AlMe_3 .

BuLi is also suitable but is used at lower temperatures and requires two equivalents of the base and two equivalents of the amine.

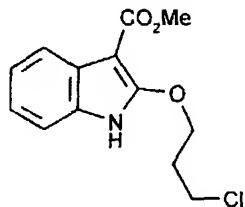
The mechanism of the reaction and role of the aluminium or lithium based catalyst is discussed in the references listed below:

Use of AlMe₃: Anwer Basha, Michael Lipton and Steven M. Weinreb,
Tetrahedron Letters, 48, 4171, 1977.

5 Use of NaH₂Et₂Al: Tae Bo Sim and Nung Min Yoon, Synlett., 1994, 827

Use of BuLi: Kim-Wenn Yang, Joseph, G. Cannon and John G. Rose,
Tetrahedron Letters, 21, 1791, 1970

The oxazinoindole compound of the formula (A) is prepared from the
10 corresponding indole by reaction with N-chlorosuccinimide and a 3-halo-propanol,
such as 3-chloropropanol or 3-bromopropanol followed by cyclisation of the
intermediate (B) by treatment with base in a suitable solvent.



15 (B)

Suitable solvents for the cyclisation include acetone and toluene, and suitable bases include potassium carbonate, aqueous sodium hydroxide.

The use of aqueous sodium hydroxide solution, even at 80°C does not cause any significant hydrolysis of the ester.

20 In the case of toluene/aqueous sodium hydroxide, a phase transfer catalyst (eg tetrabutylammonium bromide) may be added, resulting in accelerating the reaction and allowing it to proceed at a lower temperature.

The intermediate (B) may either be used as a crude oil or isolated as a white crystalline solid and then cyclised in quantitative yield to give a solution of (A) in a 25 suitable solvent (eg toluene) for coupling with the amine.

Compounds of the formula (A) and (B) are novel and form an aspect of the invention.

The following Example illustrates the invention. The following Description illustrates the preparation of an intermediate of formula (A).

5 **Example**

i) **Preparation of N-[(1-butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]-oxazino[3,2-a]indole-10-carboxamide [SB-207266]**

Method A Toluene (85L) was azeotropically dried in an argon purged reactor, cooled to 10°C and a solution of trimethylaluminium in toluene (18.57kg, 16.7% w/w, 10 43 mole) added. To this at 20 to 24°C was added a solution of 1-*n*-butyl-4-piperidinylmethylamine (7.39kg, 99.4% pure, 42.7mole) in toluene (22L) over 43 minutes. Methyl 3,4-dihydro-2H-[1,3]-oxazino[3,2-a]indole-10-carboxylate (9.65kg, 98.9% pure, 41.3 mole) was added in one portion and the reaction heated to reflux at 112°C for 4 hours 10 minutes after which time the reaction was judged to be complete 15 by HPLC analysis. 10% Sodium hydroxide solution (52.2L), prepared from 32% w/w sodium hydroxide (24L) and water (80L), was added cautiously over 16 minutes at about 60 to 70°C. The resulting mixture was heated to 70 to 80°C and the aqueous layer separated. The toluene layer was washed with 10% sodium hydroxide (52.2L) followed twice by water (29L each wash). The toluene layer was cooled and diluted 20 with hexane fraction (133L) to crystallise the product. After cooling to about 2°C overnight the product was collected by filtration, washed on the filter with hexane (21L) and dried *in vacuo* at 40°C overnight to give SB-207266 batch 207266-HP8 (12.26kg, 94.5% pure, 75.9%).

Method B 1.6M Butyllithium in hexane (1.4ml) was added to toluene (2ml) at - 25 10°. A solution of 1-*n*-butyl-4-piperidinylmethylamine (0.38g) in toluene (3ml) was added and the mixture was stirred for 5 mins. A solution of methyl 3,4-dihydro-2H-[1,3]-oxazino[3,2-a]indole-10-carboxylate (0.23g) in hot toluene (5ml) was added and the mixture was stirred at -10° for 5 mins. The mixture was diluted to 1000ml with acetonitrile:water and relative assay of the solution showed an SB-207266 content of 30 343mg (93% yield).

Method C A mixture of methyl 2-(3-chloropropoxy)-indole-3-carboxylate (100g, 0.37mol), aqueous sodium hydroxide solution (38ml, 10.8M, 0.41mol), water (38ml) and tetrabutylammonium bromide (6.0g, 0.019mol) in toluene (1000ml) was stirred at 50 - 60°C for about one hour. Water (120ml) was added and the aqueous layer was removed. The organic layer was washed with water (120ml) and dried by azeotropic distillation of toluene (250ml) giving a dry solution of methyl 3,4-dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxylate in toluene. This solution was cooled to ambient temperature and treated sequentially with a solution of 1-butyl-4-piperidinylmethylamine (66.8g, 0.39mol) in toluene (200ml) followed by a solution of trimethylaluminium in toluene (196ml, 2.0M, 0.39mol). The mixture was heated to reflux and stirred for three hours. The reaction was quenched by cautious addition of aqueous sodium hydroxide solution (460ml, 10% w/v) and then washed once with aqueous sodium hydroxide solution (460ml, 10% w/v) and twice with water (275ml each wash) while maintaining the temperature at about 70°C. Toluene (200ml) was added and the resulting solution was dried by azeotropic distillation of toluene (200ml) at about 55°C under reduced pressure. Hexane (1400ml) was added and the resulting slurry cooled to about 0 - 5°C for about one hour. The solid was isolated by filtration and dried *in vacuo* to give the product, N-[(1-butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxamide, (114.7g, 83%) as a white crystalline solid.

ii) Preparation of N-[(1-butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxamide hydrochloride [SB-207266-A]

Method A N-[(1-Butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxamide (SB-207266) (12.26kg 94.5% pure, 31.35 moles) was dissolved in acetone (70.5L) at 41°C. Anhydrous HCl in propan-2-ol (8.98L, 3.86 molar, 34.7 moles), made by dissolving HCl gas (3.1kg) in propan-2-ol (20L), was added over 8 minutes allowing the temperature to rise to 57°C. The mixture was cooled to 4°C and stirred at 2 to 4°C for 2 hours. The precipitate was filtered off, washed with cold acetone (25L) and dried at atmospheric pressure at 40 to 50°C for 17 hours to give the crude product as a white solid (12.94kg; 96.1%).

The crude product (12.94kg) was dissolved in hot ethanol (107L) and filtered through celite, washing the filter bed with further hot ethanol (18L). The filtrate was heated to 75°C and hot filtered hexane (68L) was added. The mixture was cooled to 19°C over about 4 hours and then to 4°C and stirred overnight at 1°C. The white solid was 5 filtered off, washed with a 1:1 mixture of cold ethanol/hexane (27L) and dried *in vacuo* at 50°C for 23 hours to give SB-207266A (12.36kg, 96.2% from SB-207266). This was milled in an Apex Comminuting mill through a 0.125 inch x 0.125 inch square mesh at medium speed and with hammers forward. 12.3kg (95.8% from SB-207266) was isolated as a fine, homogeneous white powder.

10 **Method B** N-[(1-Butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxamide (SB-207266) (100g, 0.27mol) was dissolved in ethanol (870ml) and the resulting solution filtered to remove particulates. Anhydrous HCl in ethanol (83ml, 3.6M, 0.30mol) was added causing the product to precipitate out of solution. The slurry was heated to redissolve the solid and hexane (550ml) was 15 added. After cooling to room temperature, the mixture was cooled to 0 - 5°C and stirred at that temperature for about two hours. The solid was isolated by filtration and dried *in vacuo* at about 40°C to give the product, N-[(1-butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxamide hydrochloride, (102.8g) in 94% yield.

20

Description

Preparation of methyl-3,4 dihydro-2H-[1,3]-oxazino[3,2-*a*]indole-10-carboxylate

Method A A solution of 3-chloropropanol (14.74kg, 98.4% pure, 153.4 mole) in dichloromethane (67L) was cooled to -17°C. In a second vessel dichloromethane 25 (68L), methyl indole-3-carboxylate (13.5kg, 99.8% pure, 76.9 mole) and 1,4-diazabicyclo[2.2.2]octane (4.75kg, assumed 100% pure, 42.3 mole) were cooled to 0°C. N-Chlorosuccinimide (11.3kg, 99.5% pure, 84.2 mole) was added to the second vessel and stirred at 0°C for 10 minutes. In the meantime methane sulphonic acid (0.59L, 99.7% pure, 6.14 mole) was added to the first vessel. The solution in the 30 second vessel was added to the first vessel over 49 minutes at -15 to 3°C and the resulting mixture stirred for a further 31 minutes at -5 to 0°C. 10% Sodium carbonate solution (147L), made from sodium carbonate (42.2kg, 398 mole) and process water

(422L), was added over 14 minutes and stirred. The organic layer was separated and washed twice with 10% sodium carbonate solution (147L each wash). The organic layer was dried over anhydrous sodium sulphate, filtered and concentrated under reduced pressure below 30°C. The concentrate was dissolved in acetone (101L) at 5 about 18°C and potassium carbonate (14.9kg) added. The mixture was stirred at 18 to 28°C for 18 hours. Analysis of the reaction showed it to be complete. The inorganic salts were filtered off, the filtrate concentrated under reduced pressure below 30°C and dissolved in dichloromethane (101L). The dichloromethane solution was washed twice with 5% sodium bicarbonate solution (85L each wash), made from sodium 10 bicarbonate (8.3kg) and water (167L) and dried over anhydrous sodium sulphate. After filtration the filtrate was concentrated under reduced pressure to a base temperature of about 95°C and diluted with toluene (12L). The toluene solution was cooled, causing the product to crystallise and cooling continued to about 0°C overnight. The product was collected by filtration, washed on the filter with cold 15 (0°C) toluene (7L) and dried *in vacuo* at 30°C for 21 hours to give the title compound (9.654kg, 98.9% pure, 53.7%).

Method B A solution of 3-chloropropanol (142.47g, 1.51 mole) in dichloromethane (1200ml) was cooled to -20°C. In a second vessel dichloromethane (1300ml), methyl indole-3-carboxylate (240.0g, 1.37 mole) and 1,4-diazabicyclo[2.2.2]octane (84.52g 0.75 mole) were cooled to 0°C. 20 N-Chlorosuccinimide (201.22g 1.51 mole) was added to the second vessel and stirred at 0°C for 10 minutes. In the meantime methane sulphonic acid (10.56ml) was added to the first vessel. The solution in the second vessel was added to the first vessel keeping the temperature below about 0°C, and the resulting mixture stirred for a 25 further 2.5 hours at -5 to 0°C. 10% Sodium carbonate solution (2500ml) was added and stirred. The organic layer was separated and washed twice with 10% sodium carbonate solution (2500ml each wash). The organic layer was dried over anhydrous sodium sulphate, filtered and concentrated under reduced pressure. The concentrate was triturated with ethyl acetate (120ml) and the mixture stirred at 0°C for about 1 30 hour. The resulting solid was filtered, washed a small quantity of ethyl acetate and dried under vacuum to give *methyl 2-(3-chloropropoxy)-indole-3-carboxylate* (202.5g) as a white crystalline solid in 55% yield.

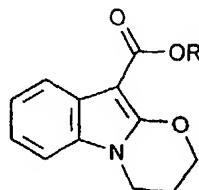
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A mixture of methyl 2-(3-chloropropoxy)-indole-3-carboxylate (81.5g, 0.304 mole), aqueous sodium hydroxide solution (31ml, 10.8M, 0.335 mole), water (31ml) and tetrabutylammonium bromide (4.9g, 0.015 mole) in toluene (815ml) was stirred at 50 - 60°C for about 45 minutes. The aqueous layer was removed and the organic 5 layer washed twice with water (100ml each wash). The resulting toluene solution was dried by azeotropic distillation of solvent (265ml) under reduced pressure (60°C, 160mbar) giving a dried solution of the title compound in toluene.

Method C A solution of 3-chloropropanol (142.47g, 1.51 mol) in dichloromethane (1200ml) was cooled to -20°C. In a second vessel dichloromethane (1300ml), methyl 10 indole-3-carboxylate (240.0g, 1.37 mol) and 1,4-diazabicyclo[2.2.2]octane (84.52g 0.75 mol) were cooled to 0°C. N-Chlorosuccinimide (201.22g 1.51 mole) was added to the second vessel and stirred at 0°C for 10 minutes. In the meantime methane sulphonic acid (10.56ml) was added to the first vessel. The solution in the second vessel was added to the first vessel keeping the temperature below about 0°C, and the 15 resulting mixture stirred for a further 2.5 hours at -5 to 0°C. 10% Sodium carbonate solution (1250ml) was added and the mixture stirred for about 30 minutes. The organic layer was separated and washed once more with 10% sodium carbonate solution (1250ml) and once with water (1250ml). The organic layer was dried over anhydrous sodium sulphate, filtered and concentrated under reduced pressure. The 20 concentrate was triturated with toluene (400ml) and the mixture stirred at 0°C for about 1 hour. The resulting solid was filtered, washed with toluene and dried *in vacuo* to give methyl 2-(3-chloropropoxy)-indole-3-carboxylate (245.5g) as a white crystalline solid in 67% yield.

Claims

1. A process for the preparation of SB 207266 or a pharmaceutically acceptable salt thereof, which process comprises the reaction of N-(1-nbutyl-4-piperidyl)methylamine with a compound of formula (A):



(A)

wherein R is alkyl.

10

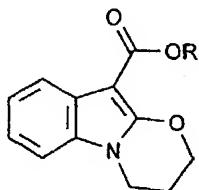
2. A process according to claim 1 wherein R is methyl or ethyl.

3. A process according to claim 1 or 2 wherein the reaction is catalysed by a aluminium or lithium based catalyst.

15

4. A process according to claim 3 wherein the catalyst is trimethylaluminium.

5. A compound of formula (A):



(A)

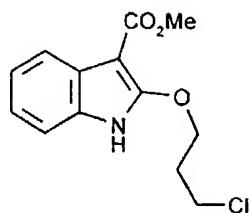
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wherein R is alkyl.

6. Methyl-3,4 dihydro-2*H*-[1,3]-oxazino[3,2-*a*]indole-10-carboxylate.

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7. A compound of formula (B):



(B)

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 97/04413

| | | |
|--|---|--|
| A. CLASSIFICATION OF SUBJECT MATTER IPC 6 C07D498/04 C07D209/42 // (C07D498/04, 265:00, 209:00) | | |
| According to International Patent Classification (IPC) or to both national classification and IPC | | |
| B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 6 C07D | | |
| Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched | | |
| Electronic data base consulted during the international search (name of data base and, where practical, search terms used) | | |
| C. DOCUMENTS CONSIDERED TO BE RELEVANT | | |
| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
| Y | <p>GASTER L M ET AL: "N-[(1-Butyl-4-piperidinyl)methyl]-3,4-dihydro-2H-[1,3]oxazino[3,2-a]indole-10-carboxamide hydrochloride: the first potent and selective 5-HT4 receptor antagonist amide with oral activity." J. MED. CHEM. (JMCMAR, 00222623);95; VOL.38 (24); PP.4760-3, SMITHKLINE BEECHAM PHARMACEUTICALS; NEW FRONTIERS SCIENCE PARK; HARLOW ESSEX; CM19 5AW; UK (GB), XP002049018 see Scheme 1 and compounds 10-12</p> <p>---</p> <p>WO 93 18036 A (SMITHKLINE BEECHAM PLC; UK) 16 September 1993 cited in the application see the whole document, especially example 3.</p> <p>---</p> <p>-/-</p> | 1-7 |
| Y | | 1-7 |
| <input checked="" type="checkbox"/> Further documents are listed in continuation of box C. | | <input checked="" type="checkbox"/> Patent family members are listed in annex. |
| <p>* Special categories of cited documents :</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"S" document member of the same patent family</p> | | |
| 1 | Date of the actual completion of the international search 3 December 1997 | Date of mailing of the international search report 16.12.97 |
| Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 apo nl, Fax: (+31-70) 340-3016 | | Authorized officer Scruton-Evans, I |

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 97/04413

| C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT | | |
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| A | WYMAN P A ET AL: "Azabicyclic Indole Esters as Potent 5-HT4 Receptor Antagonists" BIOORG. MED. CHEM. (BMECEP,09680896);96; VOL.4 (2); PP.255-61, SMITHKLINE BEECHAM PHARMACEUTICALS;ESSEX; CM19 5AW; UK (GB), XP002049019 see the whole document ----- | 1-7 |

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